ANTISECRETORY FACTOR PEPTIDES REGULATING PATHOLOGICAL PERMEABILITY CHANGES

[0001] This application is a continuation of U.S. Application No. 09/029,333, filed on March 13, 1998, which was a national stage filing under 35 U.S.C. § 371 of International Application No. PCT/SE96/01049 filed on August 23, 1996, which International Application was published by the International Bureau in English on March 6, 1997, and which claims priority to Swedish Application No. 9502936-9 filed August 24, 1995.

[0002] The present invention relates to new antisecretory factors having fluid transport and/or inflammatory reactions regulating properties as well as polynucleic regulating properties, and polynucleic acids coding therefor, and the use thereof.

[0003] All cells and tissues of the body are critically dependent on a constant and normal fluid environment in combination with an adequate blood supply. Derangement of one or both of these supporting systems may rapidly become fatal. Concerning fluid imbalance, two principally different systems exist:

A. edema, which is characterized by the abnormal accumulation of fluid in the intercellular tissue spaces or body cavities, or

B. dehydration, which, in a strict sense, means loss of water only, but is in fact commonly used to describe the combined loss of water and ions.

[0004] The most common forms of either edema or dehydration are:

diarrheas, inflammatory bowel diseases, brain edema, asthma, rhinitis, conjunctivitis, arthritis, glaucoma, various forms of pathological intracranial pressure (increase or decrease), pressure alteration in the middle ear such as Morbus Ménière,

dermatitis, chemical or physical derangement of the skin and skin adjacent glands such as mastitis, various forms of endocrine disorders, such as diabetes insipidus. Conn's syndrome, Cushing's syndrome and Morbus Addison, kidney diseases such as pyslonephritis and glomerulonephritis, metabolic diseases such as myxedema and acute intermittent porphyria, side effects during treatment with various drugs such as antidiabetics, tricyclic antidepressants, cytostatics, barbiturates, narcotics and narcotic analogues.

[0005] Diarrhea is caused by a change in the permeability in the gut for electrolytes and water. This disturbance is often caused by bacterial enterotoxins such as those produced by *Escherichia coli, Campylobacter jejuni, Vibrio cholerae, Shigella dysenteriae* and *Chostridium difficile*. The disturbance could also be caused by intestinal inflammation. Since the uptake of water is coupled to the uptake of electrolytes and nutrients, animals with frequent diarrhea suffers from malnutrition, resulting in retardation of the daily weight gain in the growing animal. The body counteracts these reactions by neuro-hormonal mechanisms such as the release of somatostatin and opiate peptides from interneurons in the intestinal mucosa. These polypeptides are capable of reversing fluid secretion and diarrhea.

[0006] The recently described antisecretory factor (AF) has been partially purified from pig pituitary gland and shown to reverse pathological secretion induced by various enterotoxins. High levels of AF in sow milk protect the suckling piglets against neonatal diarrhea.

[0007] Antimicrobial drugs have been widely used in the treatment of diarrhea in both human and veterinarian medicine. They are also used as feed additives for pigs, calves and chicken. However, due to the rapid development of resistant bacteria in the gut,

the use of antibiotics against enteritis is generally not accepted in human medicine and their use is also diminishing in veterinarian medicine.

[0008] Other antidiarrheal drugs counteract the secretion in the intestinal mucosa. Since these drugs are directed against the host animal, it is unlikely that resistance against the drugs will develop. These types of drugs include nerve-active drugs like phenothiazines and thioxanthenes. Due to some serious side effects these types of drugs have not been accepted for treatment of diarrhea in most countries. Other drugs are derivatives of opiates like codeine and loperamide since these drugs mainly acts by inhibiting intestinal mobility, they also inhibit the clearance of pathogenic bacteria from the gut and should definitely not be recommended against dysenteric bacteria or parasites. Derivatives of somatostatin have been introduced recently, but have so far a limited use due to difficulties in the administration of the drugs and possible interactions with the endocrine regulation of growth.

[0009] The antisecretory factor (AF) has so far not been used directly for treatment of diarrhea or malnutrition due to the difficulties involved in obtaining a pure preparation of this protein. However, it has been possible to induce similar proteins in domestic animals which have been given a specific feed (SE Patent No. 9000028-2). Pigs given this feed obtained high levels of AF-like proteins and had a significant increase in the daily growth rate compared to matched controls. AF in rats challenged with toxin A from *C. difficile* protects not only against intestinal secretion but also against inflammation and bleeding in the gut.

[0010] A major object of the present invention is to provide a new recombinant protein and homologues and fragments (peptides) thereof for use in normalizing

pathological fluid transport. These proteins and peptides are collectively called antisecretory factors (AF). The use of AF also partly inhibits, or totally eliminates the development of inflammatory reactions of various aetiologies. Reconstitution back to normal (fluid transport or inflammation) is obtained by the use of proteins or peptides. Further the AF proteins or peptides are effectively absorbed via various mucus membranes without losing in potency (when compared to intravenous administration). Consequently, a multitude of treatment regimens exist, and a correctly administrated protein or peptide make it possible to rapidly reconstitute a deranged fluid (water and ion) balance, an inflammatory reaction, or both.

[0011] In summary, the recombinant AF (rAF) and the homologues and fragments thereof could be used for immunodetection, as feed additive for growing animals and as antidiarrheal and drugs against diseases involving edema, dehydration and/or inflammation.

[0012] The objects of the present invention are the following:

A recombinant protein having essentially the amino acid sequence shown in SEQ ID NO:2, or homologues or fragments thereof.

[0013] A fragment of the recombinant protein shown in SEQ ID NO:2: which fragment is chosen from the group comprising

a) amino acids nos. 35-42

b) amino acids nos. 35-46

c) amino acids nos. 36-51

d) amino acids nos. 36-80

e) amino acids nos. 1-80

of the amino acid sequence shown in SEQ ID NO:2.

[0014] A peptide $X_1VCX_2X_3KX_4R$ corresponding to the fragment comprising the amino acids no. 35-42 of the recombinant protein shown in SEQ ID NO:2, wherein X is I or none, X_2 is H, R or K, X_3 is S, L or another neutral amino acid and X_4 is T or A.

[0015] Antibodies against a recombinant protein having essentially the amino acid sequence shown in SEQ ID NO:2, or homologues or fragments thereof.

[0016] A protein binding to antibodies specific to a recombinant protein having essentially the amino acid sequence shown in SEQ ID No. 1, or homologues or fragments thereof.

[0017] A composition for normalizing pathological fluid transport and/or inflammatory reactions comprising as an active principal an effective amount of the recombinant protein having essentially the amino acid sequence shown in SEQ ID NO:2, or homologues or fragments thereof.

[0018] Use of a recombinant protein having essentially the amino acid sequence shown in SEQ ID NO:2, homologues or fragments thereof for manufacturing a composition for normalizing pathological fluid transport and/or inflammatory reactions.

[0019] Feed for normalizing pathological fluid transport and/or inflammatory reactions in vertebrates, comprising as an active agent a recombinant protein having essentially the amino acid sequence shown in SEQ ID NO:2, or homologues or fragments thereof, or an organism capable of producing such a protein or homologues or fragments thereof.

[0020] A process of normalizing pathological fluid transport and/or inflammatory reactions in vertebrates, comprising administering to the vertebrate an effective amount of a recombinant protein having essentially the amino acid sequence shown in SEQ ID NO:1, or

homologues or fragments thereof, or an organism producing said protein or homologues or fragments.

[0021] Use of specific antibodies against a recombinant protein having essentially the amino acid sequence shown in SEQ ID NO:2, or homologues or fragments thereof, for detecting said protein or fragments in organisms.

[0022] Nucleic acids coding for a recombinant protein having essentially the sequence shown in SEQ ID NO:2, or homologues or fragments thereof.

[0023] Use of nucleic acids coding for a recombinant protein having essentially the amino acid sequence shown in SEQ ID No. 1, or homologues or fragments thereof, for producing corresponding proteins or homologues or fragments.

[0024] Use of probes or primers derived from nucleic acids coding for a recombinant protein having essentially the sequence shown in SEQ. ID No. 1, or homologues or fragments thereof, for detecting the presence of nucleic acids in organisms.

[0025] Vector comprising nucleic acids coding for a recombinant protein having essentially the amino acid sequence shown in SEQ. ID No. 1, or homologues or fragments thereof.

[0026] Host except human comprising a vector including nucleic acids coding for a recombinant protein having essentially the amino acid sequence shown in SEQ. ID No. 1, or homologues or fragments thereof.

[0027] A strain of an organism except human capable of producing a protein having essentially the amino acid sequence shown in SEQ ID No. 1, or homologues or fragments thereof.

[0028] As organisms capable of producing the recombinant protein use can be made of different types of organisms, such as recombinant bacteria and eucaryotic organisms, such as yeast, plants and vertebrates except humans.

[0029] Despite ten years of attempts to purify AF by conventional biochemical techniques, it has not been possible to obtain AF in a homogeneous form. However, by means of a new procedure of preparing a semipure AF for immunization and selecting antiserum by means of an immunohistochemical method a suitable antiserum was chosen. With this antiserum it has now been possible to clone recombinant human cDNA expressing AF in *E. coli*.

[0030] The sequence of the new cDNA was determined and shown to be unique. By knowledge of this sequence, oligonucleotide probes were constructed which hybridize with human and porcine pituitary RNA. The size of this RNA, about 1400 basepairs, complies with the size of the sequenced cDNA comprising 1309 basepairs plus a poly(A)tail. A partial cDNA sequence from rat pituitary gland has been shown to be identical with that of the human cDNA reflecting a ubiquitous structure conserved in AF genes from different species. This resemblance makes it possible to use the same oligonucleotide probes to identify AF-coding RNA and DNA from different species.

[0031] It has furthermore been possible to express the rAF in a biological active form. The AF protein in form of a fusion protein with gluthathione S-transferase was expressed in large amounts in *E. coli*. and purified to homogeneity by affinity chromatography. After cleavage of the fusion protein with thrombin, the recombinant AF (rAF) was shown to be extremely potent, 44 ng (10^{-12} mol) giving a half-maximal inhibition of cholera toxin-induced fluid secretion in rat intestine.

[0032] By gene technique smaller fragments of rAF was produced. The activity was shown to reside in a small sequence consisting of 7 to 8 amino acids. This was confirmed by help of chemical solid phase synthesis by which technique an octapeptide was produced and shown to be almost as biological potent as rAF on molar basis. With help of site directed synthesis a variety of sequences within the active site was constructed and replacements of certain amino acids shown to be possible without abolishing the biological activity.

[0033] The fluid secretion was measured by the intestinal loop model: a section (loop) of the small intestine is ligated by means of two sutures; in the loop a certain amount of enterotoxin is injected. If antisecretory drugs are tested they are injected between one hour before and two hours after toxin challenge. The injection was made by three different routes; intravenously, intraintestinally and intranasally. The fluid is accumulating in the loop 5 h after toxin challenge. The secretion is calculated from the weight of the accumulated fluid per cm intestine.

[0034] The sequence of the protein was determined both directly by amino acid sequencing and indirectly by deduction from the cDNA sequence.

[0035] Recombinant AF seems to exert very little toxic or systemic effects since no obvious toxic reactions were noted in rats given 100 fold higher doses than that causing half-maximal inhibition. Since it is efficient when injected in the small intestine it could be administrated perorally.

[0036] The recombinant AF inhibits secretion also when injected after toxin challenge in contrast to the preparations of natural AF tested which seem to efficient only

when injected before the toxin. Thus, rAF could be used both prophylactically and therapeutically.

[0037] Further, rAF and its peptide fragments were shown to inhibit cytotoxic reactions and inflammation in the gut caused by toxin A from *Clostridium difficile*. By help of a dye permeability test rAF and its fragments were shown to reverse pathological permeability changes induced by cholera toxin not only in the intestinal mucosa but also in plexus choroideus which regulates the fluid pressure in the brain.

[0038] Antisera against rAF were produced in rabbits and used in enzyme-linked immuno assays (ELISA). This assay might be used to measure AF in body fluids or feed.

[0039] A method of purifying antibodies against AF (natural or recombinant) by means of affinity chromatography on columns with agarose coupled rAF is reported below.

[0040] The antibodies were also shown to be efficient for detection of AF in tissue sections by means of immunohistochemical techniques and for detection of AF in Western-blot.

[0041] The invention will now be described further by means of the following non-limiting Examples together with the accompanying drawings.

Example 1. Antibodies against AF produced for cloning of cDNA

[0042] Antisecretory factor was prepared from pig blood by means of affinity chromatography on agarose and isoelectric focusing. To one liter of pig blood (containing anticoagulating substances) 1 g of sodium thiosulfate and 1 mg of phenylmethylsulfonylfluoride were added. The blood cells were separated by centrifugation and the clear plasma was eluted through a column with Sepharose 6B

(Pharmacia LKB Biotechnology Stockholm), the gel volume corresponding to about 10% of the volume of the solution. After washing with three bed volumes of phosphate buffered saline (PBS = 0.15 M NaCl, 0.05 M sodium phosphate, pH 7.2), the column was eluted with two bed volumes of 1 M α-methyl-D-glucoside dissolved in PBS. The eluate was concentrated and dialyzed against water on an "Omega 10k flow through" ultrafilter (Filtran Technology Corp.). The fraction was subsequently fractionated by isoelectric focusing in an ampholine (Pharmacia) gradient pH 4-6 on a 400 ml isoelectrofocusing column (LKB, Sweden). A fraction having an isoelectric point-between 4.7 and 4.9 was collected and dialyzed against PBS. Thus, partially purified AF was divided into small aliquotes and used for production of antiserum in rabbits according to a previously described method.

[0043] The rabbits were immunized and the sera tested for their capacity to stain intracellular material in sections of human pituitary gland (method described in Example 6). Only one of the sera showed specific and distinct intracellular staining without staining extracellular matrix proteins. This antiserum was selected for screening of a cDNA/lambda phage GT11 library from human pituitary gland expressing proteins in *E. coli*.

Example 2. Screening cDNA libraries from human pituitary gland and brain.

[0044] A 5'-stretch cDNA library from normal human pituitary gland, derived from tissues obtained from a pool of nine Caucasians, was purchased from Clontech Laboratories. For screening of the library, phages were plated at 3 x 10^4 plaque forming units per 150 mm dish on *E. coli* Y1090. The previously described rabbit antiserum against porcine AF was absorbed with 0.5 volumes of *E. coli* Y1090-lysate for 4 hours at

23°C and diluted to a ratio of 1:400 and screening performed according to Young and Davis (1).

[0045] Alkaline-phosphatase-conjugated goat anti-rabbit antibodies were used as second antibodies (Jackson). Positive plaques were picked, eluted into phage suspension medium [20 mM Tris-HC1 (pH 7.5), 100 mM NaCl, 10 mM $MgSO_4$, 2% gelatin], replated, and screened until all plagues tested were positive.

[0046] *cDNA-recloning* - Phage DNA from AF recombinants was isolated with Wizard Lambda Preps (Promega) and digested with *Eco*R1. The inserts were purified with Sephaglas BandPrep Kits (Pharmacia), recloned into pGex-lλT vector (Pharmacia) as described by the manufacturer and transfected into Epicurian Coli XLI-Blue, Top 1 cells or BL21 cells (all three from Stratagen). rAF or rpeptides were prepared in BL21 cells when not stated otherwise (2).

[0047] *Amplification of cDNA by PCR* - To obtain the missing 5'-end of the cDNA a PCR-based method called RACE (rapid amplification of cDNA ends) was performed. A modified RACE-method that generates 5'-RACE-Ready cDNA with an anchor oligonucleotide ligated to the 3'-ends of the human brain cDNA molecules was purchased from Clontech Laboratories. The 5'-end was amplified from a portion of the 5'-RACE-Ready cDNA in two PCR amplification steps using a 5' primer complementary to the anchor and two nested gene-specific 3' PCR primers A and B (A=base 429-411 and B=base 376-359; Fig. 1a). Various smaller portions of the RACE fragment was further amplified in order to express the corresponding peptides and test for their biological properties. The position of the base and amino acid at the start and end of these oligonucleotide fragments and their corresponding peptides are shown in Table 1. Porcine

and bovine cDNA (Clontech Laboratories) was used as templet for amplifying fragments corresponding to N3 in Table 1. Variation of the sequence was also inserted artificially by site directed mutagenesis in which method various oligonucleotides corresponding to position 168-193 was synthesized in order to replace one by one of amino acid 35-42 (positions as shown in SEQ ID NO:2). The amplified DNA fragment was cloned into pGex-1λT vector by using the EcoR1 site built into the anchor and the gene-specific primer. To verily the sequence obtained by the RACE method, double stranded cDNA from human pituitary gland and brain (Clontach) were amplified with primer pair C/D containing an extra EcoR1-cleavage site (Fig. 1b). The primers were designed to allow the entire open reading frame (ORF) to be amplified. The pituitary and brain PCR-products of expected size were digested with ECOR1, isolated and cloned into the plasmid pGex-1λT vector.

[0048] *DNA sequencing and oligonucleotides* - DNA from plasmid pGex-1λT was used as a template for sequencing of the inserts by dideoxy-chain-termination method (15) using the Sequence version 2.0 kit (U.S. Biochemical Corp.). Initial forward and reverse primers copying regions of pGex-lλT immediately upstream and downstream of inserted DNA were obtained from Pharmacia. Subsequent primers were synthesized (Scandinavian Gene Synthesis AB) on the basis of sequence information obtained. Three different PCR clones were sequenced in order to avoid base-exchange by *Taq* polymerase in the 5'-RACE method.

[0049] Nucleotide sequence and the deduced protein sequence data were compiled and analyzed by using MacVector 4.1 (Eastman Chemical Co.). To predict the corresponding amino acid sequence of the cDNA inserts, codon usage of different reading

frames was compared and gave one large open reading frame. Interrogation of DNA and protein sequence data was carried out by use of an Entrez CD-ROM disc (National Center for Biotechnology Information, Bethesda, USA).

[0050] *Molecular cloning and sequence analysis of cDNA* - Polyvalent antisera against AF protein from pig were used for screening cDNA from human pituitary glands. Two clones expressing immunoreactive AF were isolated, rescued from phage lambda and recloned into the EcoR1 site of vector pGex-1λT as described in the kit provided from Pharmacia. Restriction analysis gave insert sizes of 1100 and 900 bp, respectively DNA-sequencing of the two clones revealed homology to be complete except for one substitution (Fig. 1, C replacing T at position 1011). A sequence upstream of the 5'-end of clone 2 was obtained by means of the RACE method. The fragment had a total length of 376 bp (not including the synthetic nucleotide arm at the 5'-end). The total reconstructed cDNA contained 1309 basepairs followed a poly-A tail, which was preceded by a poly-A signal (Fig. 1, positions 1289-1295). An open reading frame (ORF) of 1146 bp (positions 63-1208) was identified.

Example 3. Expression of mammalian AF protein from recombinant plasmids.

[0051] *Construction and purification of fusion proteins* - The cDNA-clones obtained by immunological screening and by PCR amplification of the entire cDNA were ligated to pGex-1λT. This vector allows expression of foreign proteins in *E.coli* as fusions to the C terminus of the *Schistosoma japonicum* 26 kDa glutathione S-transferase (GST), which can be affinity purified under nondenaturing conditions with help of the kit provided from Pharmacia. Briefly, overnight cultures of *E.coli* transformed with recombinant pGex-1λT

plasmids were diluted in fresh medium and grown for a further 3 h at 37°C. Protein expression was induced by 0.1 mM IPTG (isopropyl-beta-D-thiogalactopyranoside), and after a further 4 h of growth at 30°C, the calls were pelleted and resuspended in PBS. Cells were lyzed by sonication, treated with 1% Triton X-100 and centrifuged at 12000X g for 10 min; the supernatant containing the expressed fusion proteins was purified by passing the lysates through glutathione agarose (Pharmacia). The fusion proteins were either eluted by competition with free glutathione or were cleaved overnight with 10 U bovine thrombin to remove the AF-protein from the GST affinity tail. The entire method of using the pGex plasmid and purifying the recombinant proteins or peptides was performed by means of the kits provided from Pharmacia.

[0052] *Sequence and size of Recombinant AF-proteins* - To confirm the coding sequence, the full-length transcript was isolated by using PCR-amplification of pituitary and brain cDNA. Using the primer pair C/D, 1215 bp identical to the sequence of clone-4 (Fig. 1, SEQ ID NO:1) was isolated. The open-reading frame encoded 382 amino acids with a calculated molecular mass of 41.14 kDa and a calculated pI of 4.9.

[0053] The AF clones-1, 2 and 3 as well as the oligonucleotides N1-N5 (Fig. 1 and Table 1) were ligated into the pGEX-1λT plasmid vector so that the ORF was in frame with the glutathione S-transferase (GST) protein. The constructs were transformed into *E.coli*, and expression of fusion proteins was induced with IPTG. The purified fusion proteins and the thrombin-cleaved AF protein or peptide were subjected to SDS-PACE and Western blotting using antiserum against porcine antisecretory factor (Fig. 2). Coomassie brilliant blue staining of the proteins revealed discrete bands for each protein except for the GST-AF-1 protein which manifested degradation into smaller components.

[0054] *Solid phase peptide synthesis* - Smaller peptides (P_7 to P_{18} in Table 1) was produced (K.J. Ross-Petersen AS) on solid phase in an Applied Biosystems peptide synthesizer. The purity of each peptide was 93-100% as evaluated on reversed phase HPLC on Deltapak C18, 300 A using a linear gradient of 0.1% trifluoro acetic acid in water/acetonitril.

[0055] *Amino acid sequencing* - Protein sequence analysis was performed to further validate the identified ORF. The pure AF proteins were run in 10% macro-slab gel SDS-PAGE (14) and the proteins transferred to a Problot membrane (Applied Biosystems) by electroblatting (Bio-Rad). Spots, visualized by Ponceau S staining, were excised from the blot and the first 20 amino acids of the proteins were sequenced by automated Edman degradation on an automatic sequencer (Applied Biosystems).

[0056] The N-terminal sequences of clone-2 and clone-3 were determined, and shown to perfectly match amino acids 63-75 and 130-140, respectively, of the predicted sequence (Fig. 1, SEQ ID NO:2).

[0057] Comparison with other protein sequences available from GenBank revealed that the sequence of rAF (Fig. 1, SEQ ID NO:2) is unique in all its parts and no similar sequence has been reported.

[0058] The first ten residues of the protein appear to be relatively hydrophobic when analyzed according to KyteDoolittle (22) and might constitute a signal peptide, which is cleaved out prior to exocytosis of the protein. This interpretation is supported by the Western blot analyzes (Fig. 3) in which the recombinant protein appeared to have a slightly higher molecular mass than the protein extract from pituitary gland. Some of this

difference, however, might also be due to the additional five amino acids in the recombinant protein constituting the trombin cleavage site of the fusion protein.

Example 4. Production and testing antisera against rAF

[0059] *Antisera against recombinant GST-AF fusion protein* - Antibodies against the purified fusion proteins GST-AF-1, GST-AF-2 and thrombin-cleaved pure AF-I protein (=rAF) for use in ELISA, Western blot and immunohistochemical studies were produced in rabbits. Each rabbit was given 100 pg of antigen In I ml PBS mixed with an equal volume of Freund's complete adjuvant; each immunization was distributed in 8-10 portions injected in the back intracutaneously. Two booster dozes with 50 μg antigen were injected at 3 and 5 weeks, the last one without Freund's complete adjuvant. The rabbits were bled 6 days after last booster and sera were prepared and stored at -20°C. The sensitivity of the antiserum was tested with a dot blot assay. GST-AF-2 was applied on an ECL nitrocellulose membrane in 1/5 dilutions, and the antiserum diluted 1:1000. The membrane was blocked with 1% bovine serum albumin (BSA) in PBS at 4°C for 16 h, and then incubated for 1 ½ h with a 1:800 dilution of rabbit anti-GST-AF or porcine AF antiserum. The blot was developed with alkaline phosphatese-conjugated goat anti-rabbit immunoglobulin followed by 5-bromo-4-chloro-3-indolyl phosphate and p-nitro blue tetrazolium (Boehringer Mannheim). The estimated limit for antigen detection was about 1 ng in this test.

[0060] *SDS-polyacrylamlde gel electrophoresis and immunoblotting* - SDS-polyacrylamide gel electrophoresis (SDS-PAGE) of human and porcine pituitary gland extracts and pure AF-proteins was performed in 10% acrylamide minislab gels, essentially

as described by Laemmli (4) with the modification that bis-acrylamide as a cross-linker was replaced by *N,N'*-diallyltartardiamide with the corresponding molarity. Pyronin Y (Sigma) was used as a marker of the electrophoretic front. Prestained molecular weight reference were purchased from BDH. Proteins were then either stained with Coomassie brilliant blue or transferred electrophoretically to 0.45 mm pore-size ECL nitrocellulose (Amershem) for immunoblotting. The subsequent incubations with BSA, conjugated anti-IgG and alkaline phosphatase substrate were the same as for the dot blot assay described above.

[0061] As stated above Coomassie Brilliant Blue staining revealed no discrete band for the GST-AF-1 protein, which was probably due to proteolytic degradation into smaller components. However, in the Western blot analyzes the full length protein gave a much stronger signal than the degradated products (Fig. 2b). The strong reaction with the antiserum against porcine AF indicated that the recombinant proteins indeed have the same immunoreactivity as AF. The molecular weight of the full length protein appeared to be about 60 kDa which is higher than the true mol. wt of 41139 Da estimated from the amino acid composition. Furthermore, the proteins were also immunoblotted and probed with antiserum raised against GST-AF-2, which bound to the thrombin-cleaved proteins (Fig. 3).

[0062] Antiserum against recombinant GST-AF-2 reacted with the naturally occurring AF protein of an apparent mol mass of 60 kDa, and with some smaller components, probably enzymatic degradation products (Fig. 3 a).

[0063] *ELISA for determination of AP-concentrations* - ELISA assays were performed using anti-AF-1 and anti-AF-2 according to a previously described method (5). As shown in Fig. 3b the sensitivity of the test with the crude antiserum was between 1-10

μg protein whereas the test with the affinity purified antibody had a sensitivity between 5 and 50 ng protein,

Example 5. Northern blot analysis of RNA from pituitary gland

[0064] *Northern blot analysis* - Human pituitary glands were obtained postmortem from Sahlgrenska Hospital (permission given by Swedish Health and Welfare Board; 2% transplantationslagen, 1975:190). To obtain RNA, pituitary glands were extracted with guanidinium thiocyanate RNA according to Chomczynski and Sacchi (6). Polyadenylated RNA was selected by means of a commercial kit (Pharmacia) using columns with oligodT-cellulose. In addition, a pool of human pituitary mRNA from 107 individuals purchased from Clontach was used. Five μg of each sample of poly(A+)RNA was glyoxal-treated and electrophoresed in a 1.2% agarose gel (7). After capillary alkaline transfer for 3 h in 0.05 M NaOH to Hybond N+ nylon membranes (Amersham), prehybridization and hybridization were carried out for 24 h each at 42°C. The hybridization solution contained 50% formamide, 5xSSPE, 10XDenhard's solution with 250 μg/ml denaturated low-MW DNA and 50 μg/ml polyadenylic acid. The blots were probed with four different antisense 28 bp oligonuclaotides comprising the positions 132-105 (primer E), 297-270 (primer F), 748-721 (primer G) and 833-806 (primer H) of the sequence (Fig. 1); the probes were 3'-end labelled with terminal transferase (Boehringer Mannheim) plus [α^{32p}]ddATP (Amersham) and purified on Nick columns (Pharmacia). Five postwashes in 5XSSPE/0.1% SDS - 0.5XSSPE/0.1% SDS were made at 42°C for 30 min each time, with a repeat of the last wash. Filters were exposed to Hyperfilm MP (Amersham) for 7 days.

[0065] *Expression in pituitary gland* - Northern blot analyzes were performed with a mixture of four oligonucleotide probes hybridizing with different sequences along the cloned cDNA (Fig. 4). The probes hybridized with a single band of about 1400 bp in the separated mRNA from pituitary gland. The strongest signals were obtained with the human material, but the porcine material also cross-reacted.

Example 6. Distribution of AF in sections of pituitary gland

[0066] *Species and tissues* - Human pituitary glands were obtained postmortem from Sahlgrenska Hospital (permission given by the Swedish Health and Welfare Board; §2 transplantationslagen, 1975:190). Glands were kept frozen at -70°C, except those used for histological examination which were fixed for 24 h in 4% paraformaldehyde dissolved in phosphate-buffered saline (PBS=0.15 M NaCl, 0.05 M sodium phosphate, pH 7.2) and thereafter transferred to 7.5% sucrose in PBS. Pituitary glands from pigs, 5-7 months old, obtained from a slaughter house, were placed on dry ice during transport and kept frozen at -70°C until used. Sprague-Dawley rate, 2-3 months old, were obtained for bioassay from B & K Universal AB, Sollentuna, Sweden. Rabbits (New Zealand White) for immunizations were obtained from Lidköping Kaninfarm, Sweden.

[0067] *Immunohistochemistry* - The fixed pituitary glands were frozen in liquid nitrogen, and cryo sections, 7 μm thick, were prepared. From each sample 5-10 sections comprising different parts of the gland were fastened to microscope slides. The sections were blocked in 5% fat-free dried milk and incubated with primary rabbit antiserum (anti-GST-AF-2 fusion protein) diluted 1:4000-1:8000 in a humid chamber overnight at 4°C. After rinsing in buffer, the specimens were incubated for 1 h at 23°C with alkaline

phosphatase-conjugated swine anti-rabbit immunoglobulins diluted 1:50 (Dako A/S). The immunoreaction was visualized with phosphatase substrates as described elsewhere (8). Control sections were incubated with immune serum absorbed with an excess of OST-AF-2 protein or with all incubation steps except the primary antibody.

[0068] *Distribution of AF in sections of pituitary gland.* The distribution of AF in sections of human pituitary glands was studied with immunohistochemical techniques (Fig. 5). In all specimens investigated, a moderate number of cells in the adenohypophysis were stained; the immunostained material appeared to be located in granules in the cytoplasm; preabsorption of the immune serum with an excess of GST-AF-2 protein abolished the signal. No staining was observed in the posterior part (neurohypophysis).

[0069] The distribution of immunoreactive material in the pituitary gland demonstrated solely intracellular distribution of AF in secreting cells of the anterior lobe (adenophypophysis). The proteins emanating from this lobe include growth hormone, thyrotropin, corticotropin, prolactin and lutainising hormone. The passage of these hormones from intracellular localization to the vascular system is triggered by releasing factors produced by neuroandocrinic cells in the hypothalamus.

Example 7. Biological activity of rAF

[0070] *Antisecretory activity* - The antisecretory activity was measured in a rat intestinal loop model previously described (9). A jejunal loop was challenged with 3 μg of cholera toxin. Either different doses of purified AF-1-proteins or PBS (control) was injected before or after the challenge with cholera toxin. The weight of the accumulated

fluid in the intestinal loop (mg/cm) was recorded after five hours. Each AF preparation was tested in at least six rats. Fisher's PLSD was used for statistical analysis of the data.

[0071] *Biological activity of rAF protein* - The biological activity of the pure rAF protein of clone-1 produced in *E.coli* was tested in a rat model. The capacity of the rAF to inhibit intestinal fluid secretion when injected intravenously 20-30 sec before intestinal challenge with cholera toxin is shown in Fig. 6. In control animals injected with buffer only, the cholera toxin caused a pronounced secretion, 412 ± 9 mg fluid per cm intestine. The pure rAF caused dose-dependent inhibition of the cholera secretion which was significantly different from the response to the buffer ($p < 0.01$, $n = 6$). Nine ng of clone-1 protein is sufficient to reduce the response by 34%, whereas 44 ng (10^{-12} mol) and 220 ng reduced it by 46% and 78%, respectively. The biological activity of recombinant AF is greater than that of any enterotoxin known to us and greater than that of any intestinal hormone or neuropeptide modifying water and electrolyte transport. Moreover, the level of activity of human rAF in rat is surprisingly high which probably reflects a ubiquitous structure conserved in rAF molecules from different species. This hypothesis is supported by the cross-reactivity between human and porcine material obtained in the Western blot and Northern blot analyzes.

[0072] The capacity of 0.5 μg of rAF to inhibit intestinal secretion when injected intravenously 20-30 sec before and 90 min after cholera toxin challenge was compared (Fig. 7). Both administrations gave significant inhibition compared to control animals ($p < 0.01$, $n = 6$). Thus, in contrast to natural AF, the recombinant protein was also efficient when given after toxin challenge which make rAF useful for therapeutic treatment of diarrhea.

[0073] 3 μg rAF was injected in a 8-10 cm long loop placed immediately proximal to the loop which was challenged with cholera toxin. The rAF was either induced 20-30 sec before or 90 min after the toxin-challenge. In. Fig. 8 it is shown that both test groups obtained a significant reduction of the fluid secretion compared to controls ($p<0.01$, $n=6$); no difference was observed between the two test groups. This experiment suggests that rAF is active after oral administration and might be used as an additive in animal feed provided that no serious side effect is obtained.

[0074] In the Examples described above, the rAF was produced in Epicurian Coli XL-1 cells. In these cells much of the produced rAF was degradated into smaller peptides. When rAF was produced in BL21 cells only a small portion of the rAF was degradated while in Top 1 cells no degradation was observed. Surprisingly the biological activity was proportional to the extent of degradation, i.e. more degradation resulted in higher activity. Therefore various shorter fragments were produced in order to test for their possible biological activity.

[0075] As shown in Table 1, these fragments were tested intravenously prior to cholera toxin challenge in the same way as described above for the intact rAF. The peptides expressed by clone 2 and 3 tested in amounts of 0.1, 1 and 10 μg had no effect on the toxin response. In contrast one microgram of the peptide expressed by the RACE fragment (clone 4) had a pronounced effect. A lot of shorter constructs were made from the RACE fragment and expressed in pGex-l-lambda. As shown in Table 1, the active site was found to be situated between amino acid residue 35 to 51. In order to determine more exactly the active site three small peptides were made by solid phase peptide synthesis.

Two of them were active, peptide 35-46 (P3) and peptide 35-42 (P1); the latter octapeptide IVCHSKTP, (P1) was active in a dose less than 1 ng being almost as active in a molar basis as the intact rAF. In contrast a shorter hexapeptide VCHSKT (P2) exerted no effect when tested in doses between 1 ng and 10 μg.

[0076] A peptide $X_1VCX_2X_3KX_4R$ corresponding to the human fragment P1 but with certain changes and/or deletions, have also been produced by site directed mutagenesis and tested for biological activity. Comparison was also made of sequences from bovine and porcine cDNA. These studies suggested the following changes and/or deletions:

X_1 is I or none

X_2 is H, R or K

X_3 is S, L or another neutral amino acid

X_4 is T or A.

Table 1

Code	Oligonucleotide*	Peptide**	Inhibition of cholera secretion***	
				pmol ED50
N1	63-301	1-80	+	4
N2	168-301	36-80	+	6
N3	168-215	36-51	+	3
N4	122-170	21-36	-	
N5	186-269	42-69	-	
P3	S.P.S.****	35-46	+	7
P1	S.P.S.	35-42	+	5
P2	S.P.S.	36-41	-	

[0077] * Position of the basepair in (SEQ ID NO:1) which was expressed in the construct made from the AF cDNA and pGex-l-lambda.

[0078] ** Positions of the amino acid-residues (SEQ ID NO:2) in AF at the start and end of the synthesized peptide.

[0079] *** Inhibition of cholera toxin induced fluid secretion in a ligated rat intestinal loop; the amount (pmol) causing halfmaximal inhibition (ED50) is noted for active peptides.

[0080] **** These peptides were produced by solid phase synthesis.

[0081] The effect of rAF on inflammation in the intestinal mucosa was also tested in the rat intestinal loop model. Thus, 20 rate were challenged with 0.5 μg of toxin A from *Clostridium difficile* (10) and the inflammatory and fluid secretion measured after 2.5 and 5

hours, respectively (10 + 10 rats). Half of the rats in each group received 100 ng of rAF intravenously 30 sec prior to the challenge; the other half received PBS buffer as control. After killing the rat, the loops was dissected out, and the middle 2-3 cm part of the loops were frozen on dry ice. The frozen specimens were then sectioned in 8 μm thick sections by use of a Leica cryostat. The sections were stained to demonstrate alkaline phosphatases by enzyme histochemistry. Alkaline phosphatases are expressed by the intestinal epithelial cells and the staining allows an assessment and of the integrity of the intestinal epithelium.

[0082] The results revealed (Fig. 9) that the control rats developed extensive damage of the intestinal mucosa: after 2.5 h shedding of epithelial calls from the basal membrane was observed together with necrotic tissue, whereas extensive bleeding was observed after 5 h. In contrast, animal treated with rAF developed no shedding, necrosis or bleeding. The toxin A-induced fluid secretion was also inhibited from 199 ± 4 to 137 ± 5 mg/cm after 2.5 h ($p < 0.01$) and from 421 ± 3 to 203 ± 6 mg/cm after 6 h (5 rats/group, $p < 0.01$).

[0083] A similar experiment was performed with 0.5 μg of the peptide IVCHSKTR (=P1) replacing the rAF protein. The octapeptide achieved the same effect on toxin A-induced intestinal inflammation and fluid secretion as shown in Fig. 9.

[0084] *Toxicity* - In order to test the toxicity of rAF it was injected in a high dose, 50 μg per rat. No obvious toxic reaction was registered during an observation period of one week.

Example 8. Biological activity of rAF on intestinal permeability

[0085] In order to evaluate the effect of rAF on the permeability of an organic substance dissolved in the blood a test with Evans blue dye was performed according to a previously described method (11). The experiment was initially performed as described above in Example 7 and Fig. 5 with intravenous injection of rAF prior to cholera toxin challenge. However, no fluid secretion was measured but 90 min after toxin challenge Evans blue dye, 1 ml of a 1.5% solution in PBS, was injected intravenously. The dye was allowed to circulate for a 5 min long period. Thereafter the rat was subjected to transcardial perfusion via the left ventricle - right atrium (using a peristaltic pump [Cole Parmer Instruments, Chicago, Ill., USA]) with 200 ml of 4°C PBS/Alsevier's (1/1 ratio) solution during a period of some 150 sea, performed under ether anaesthesia. This procedure was undertaken in order to remove all of the EB present in the vascular system, leaving only the EB in the interstitial tissue to be detected by the formamide extraction of the dye.

[0086] The results in Table 2 demonstrate that CT-challenge significantly ($p < 0.001$) increases the amount of EB that can be extracted from the intestinal tissue with some 43%, while an intravenous injection of 1 BrT prior to cholera toxin challenge prevent this increase, i.e. the amount of EB extracted from the tissue in group 1 (control) did not differ from that in group 3 (1 rAF + CT)

Table 2

Group	Challenge	ng	EB/g int. tissue x 10^{-07}	% increase of EB-kono
1	PBS+PBS	6	29.3±1.0	-
2	PBS+CT	6	51.8±1.3	43 ($p<0.001$)
3	1rAF+CT	6	29.6±1.5	0 NS

[0087] The results shown in rigs 10 and 11 demonstrate the extravasation of the azo dye Evans blue in the small intestine and in the corresponding plexus choroideus from the lateral ventricles of the brain after intestinal challenge with cholera toxin, with and without previous treatment of the rats with P1 (IVCHSKTR),

[0088] The experiments were performed in the following way: Male Sprague-Dawley rats, weighing 350 g, were starved for 18 h prior to the experimental procedure, but had free excess to water. The rats were used in groups of six. The peptide P1, cholera toxin (CT), and PBS were administrated according to Table 3.

Table 3

Group	iv inj.1*	po.inj.*	iv.inj.2*
A	P1	CT	EB
B	PBS	CT	EB
C	PBS	PBS	EB

[0089] * P1(iv.) injection 1 were given in a volume of 2 ml PBS, the peroral (po.) injection were given in a volume of 5 ml, the intravenous

injection 2 consisted of 1.5 ml of 3% Evans blue dissolved in PBS. Ether was used for anaesthesia during the performance of all injections.

[0090] The i.v. injection of P1 (0.5 μg) or of PBS were performed 10-15 sec before the peroral challenge with 100 μg CT or with PBS; 60 min after the peroral challenge, the rats were anaesthetized with ether and injected iv. with Evans blue. The dye was allowed to equilibrate for another 30 min, whereafter the rats were again anaesthetized with ether and perfused intracardially via the left ventricle with 250 ml of Alsevers solution/PBS = 50/50, in order to remove all dye present in the vascular system. After this perfuming treatment, performed during some 2-3 min, the fluorescence registrated should represent dye present only outside the vascular system.

[0091] The brain and a part of the small intestine were sampled and frozen on dry ice and cryostat sections, 8 μm thick, were prepared. The sections were air-dried and mounted in a xylene-containing mounting media. The sections were viewed in a Zeiss fluorescence microscope using a filter combination identical to that used for rhodamin-emitted fluorescence.

[0092] The results in Figs 10 and 11 demonstrate that the fluorescent intensity (white color) is of a similar magnitude in both the small intestine (Fig. 10) and in the plexus choroideus (Fig. 11) in group A (Pl iv + CT po) and C(PBS iv + PBS po). Compared to the high fluorescent intensity in the small intestine as well as in the plexus choroideus in group B (PBS iv + CT po), the results clearly demonstrate that injection of the octapeptide prior to toxin challenge inhibits the CT-induced extravascular penetration of

Evans blue. The results suggest that this holds true not only in the vascular system of the small intestine, but also in the plexus choroideus of the lateral ventricles of the brain.

[0093] In conclusion: the effect of intravenous octapeptide IVCHSKTR administration inhibits cholera toxin-induced extravascular penetration of Evans blue in the small intestine as well as in the plexus choroideus in the central nervous system. Thus, the action of rAF and its peptide derivatives is not confined to the small intestine only, but influences also the permeability of blood vessels in the central nervous system. These findings indicate that rAF and its peptide derivatives can be used to reverse pathological intracranial pressure, pressure alteration in the middle ear and various forms of permeability changes in blood vessels.

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FIGURE LEGENDS

[0094] Fig. 1a and continued on Fig. lb. Nucleic acid sequence (SEQ ID NO:1) and deduced amino acid sequence (SEQ ID NO:2) of the new human protein. The confirmed amino acid sequence is underlined. Fig. 1c. Horizontal map showing cloned cDNA and oligonucleotide primers.

[0095] Fig. 2. Coomassie brilliant blue-stained SDS-polyacrylamide minlgel (A) and immunoblot probed with antisera against porcine AF (B). Lanes with unprimed numbers contain glutathione-agarose-purified GST-AF fusion proteins AF-1, AF-2 and AF-3, whereas lanes with primed numbers contain the fusion proteins cleaved with thrombin. Molecular weight references (R), (BDH), are indicated on the left. The GST-AF-1 fusion protein is highly degraded but the immunoblot analysis shows only the detection of a full-length protein and spontaneous thrombin cleavage product. There is a 26 kDa product in the GST-AF-3 protein, probably the glutathione S-transferase-tail that has been independently expressed.

[0096] Fig. 3a. Western blot using antiserum against recombinant protein AF-2. To the left, porcine (P) and three human (H1, H2, H3) pituitary glands; and to the right, the three recombinant proteins AF-1, AF-2 and AF-3 (see Fig. 2) were applied; in the center the molecular weight standard (R).

[0097] Fig. 3b. Enzyme linked immuno-assay (ELISA) of rAF using crude antiserum and affinity purified antibodies raised in rabbit.

[0098] Fig. 4. Autoradiogram of Northern blots of RNA from a human and porcine pituitary gland (p = pooled and i = individual material). Five μg of purified mRNA was

applied in each basin; 3'-end ^{32}P-labelled oligonucleotide probes were used and the autoradiogram developed after 7 days,

[0099] Fig. 5 Cryosections of adenohypophysis stained with antiserum against recombinant protein GST-AF-2. A. Sections incubated with immune serum showing scattered cells with varying degrees of positive immunoreactivity (solid arrows). Many calls completely lack staining (open arrows). B. Serial sections to A incubated with immune serum preabsorbed with excess of recombinant protein GSTAF-2, There is no specific staining of the cells. C and D. Larger magnifications of immunopositive cells demonstrating cytoplasmatic staining of the endocrine cells, n = nucleus, c = cytoplasms.

[00100] Fig. 6. Biological activity of recombinant protein AF-1 testing inhibition of cholera toxin-induced fluid secretion. Graded doses of the protein were injected intravenously in rat; three μg of cholera toxin was injected into an intestinal loop; after five hours the accumulated fluid (mg/cm intestine) in the loop was measured. Each value represents the mean $\pm$ S.A.E. of a group of six animals.

[00101] Fig. 7. Biological activity of intravenously injected rAF-1; 0.5 μg of rAF was administrated 20-30 sec before or 90 min after challenge with 3 μg of cholera toxin in an intestinal loop of rat.

[00102] Fig. 8. Biological activity of intraluminarly injected rAF-1; 3 μg of rAF was injected 20-30 sec before or 90 min after challenge with 3 μg of cholera toxin in an intestinal loop of rat; the rAF was injected about 5 cm proximate to the loop in which the toxin was injected.

[00103] Fig. 9 A (x 2.5) is control (PBS) loops showing cellular debris in the intestinal lumen (L), but no staining of the remaining mucosa, which suggests a total

destruction of the epithelial lining. B (0.5 μl of P1 prior to toxin challenge) shows a clearly delineated epithelial lining forming villi, suggesting a conserved and normal intestinal mucosa. L = intestinal lumen. Bars = 500 μm. C (x 10) shows the destructed mucosa in the PBS-treated control group, and D shows the corresponding mucosa in the experimental (Pl-treated) group. The black arrow point at the epithelial lining, LP = lamina propria, mm = muscularis mucosa, open arrow point at the crypt cells. Bars = 100 μm. E (x 25) shows the destructed mucosa in the control (PBS-treated) group, and F shows a corresponding magnification from a rat subjected to P1 treatment prior to toxin challenge. Bars = 50 μm.

[00104] Fig. 10. Evans blue fluorescence in jejunal specimens from three groups of rats treated with cholera toxin (CT) or control buffer (PBS); pretreatment with antisecretory peptide P1 or control buffer (PBS). LP = lamina propria. Black arrow indicating epithelial cell lining; open arrow head indicating crypt cells. Bars = 100 μm.

[00105] Fig. 11. Evans blue fluorescence in plexus choroideus specimens from the rats shown in Fig. 10. Bars = 50 μm.

Sequence Listing
SEQ ID NO: 1
SEQUENCE TYPE: Nucleotide with corresponding protein
SEQUENCE LENGTH: 1309 base pairs plus poly(A) tail

STRANDEDNESS: single
TOPOLOGY: Liniar
MOLECULE TYPE: cDNA
ORIGINAL SOURCE ORGANISM: human
IMMEDIATE EXPERIMENTAL SOURCE: Pituitary gland

FEATURES: from 63 to 1208 bp mature protein
from 1289-1295 bp poly(A) signal sequence

AATTGGAGGAGTTGTTGTTAGGCCGTCCCGGAGACCCGGTCGGGAGGGAG

CAAGGTGGCAAG ATG GTG TTC GAA AGC ACT ATG GTG TGT GTG GAC AAC AGT
101
Met Val Leu Glu Ser Thr Met Val Cys Vol Asp Asn Ser>
5 10

GAG TAT ATG CGG AAT GGA GAC TTC TTA CCC ACC AGG CTG CAG GCC CAG
149
Glu Tyr Met Arg Asn Gly Asp Phc Leu Pro Thr Arg Leu Gln Ala Gln>
15 20 25

CAG GAT GCT GTC AAC ATA GTT TGT CAT TCA AAG ACC CGC AGG AAC CCT
197
Gln Asp Ala Yal Asn Ile Val Cys His Ser Lys Thr Arg Ser Asn Pro>
30 35 40 45

GAG AAC AAC GTG GGC CTT ATC ACA CTG GCT AAT GAC TGT GAA GTG CTG
245
Glu Asn Asn Val Gly Leu Ile Thr Leu Ala Asn Asp Cys Glu Val Leu>
50 55 60

ACC ACA CTC ACC CCA GAC ACT GGC CGT ATC CTG TCC AAG CTA CAT ACT
293
Thr Thr Leu Thr Pro Asp Thr Gly Arg Ile Leu Ser Lys Leu His Thr>
65 70 75

GTC CAA CCC AAG GGC AAG ATC ACC TTC TGC ACG GGC ATC CGC GTG GCC
341
Val Gln Pro Lys Gly Lys Ile Thr Phe Cys Thr Gly Ile Arg Val Ala>
80 85 90

CAT CTG GCT CTG AAG CAC CGA CAA GGC AAG AAT CAC AAG ATG CGC ATC
389
His Leu Ala Leu Lys His Arg Gln Gly Lys Asn His Lys Met Arg Ile>
95 100 105

ATT GCC TTT GTG GGA AGC CCA GTG GAG GAC AAT GAG AAG GAT CTG GTG
437
Ile Ala Phe Val Gly Ser Pro Val Glu Asp Asn Glu Lys Asp Leu Val>
110 115 120 125

AAA CTG GCT AAA CGC CTC AAG AAG GAC AAA GTA AAT GTT GAC ATT ATC
485
Lys Leu Ala Lys Arg Leu Lys Lys Glu Lys Vol Asn Val Asp Ile Ile>
130 135 140

AAT TTT GGG GAA GAG GAC GTG AAC ACA GAA AAC CTG ACA GCC TTT GTA
533
Asn Phe Gly Glu Glu Glu Val Asn Thr Glu Lys Leu Thr Ala Phe Val>
145 150 155

AAC ACG TTG AAT GCC AAA GAT GGA ACC GGT TCT CAT CTG GTG ACA GTG
581
Asn Thr Leu Asn Gly Lys Asp Gly Thr Gly Sar His Leu Val Thr Val>
160 165 170

CCT CCT GGG CCC AGT TTG GCT GAT GCT CTC ATC ACT TCT CCG ATT TTG
629
Pro Pro Gly Pro Ser Leu Alo Asp Alo Leu Ile Ser Ser Pro Ile Leu>
175 180 185

GCT GGT GAA GGT GGT GCC ATG CTG GGT CTT GGT GCC AGT GAC TTT GAA
677
Ala Gly Glu Gly Gly Ala Met Leu Gly Leu Gly Ala Ser Asp Phe Glu>
190 195 200 205

TTT GGA GTA GAT CCC ACT GCT GAT CCT GAG CTG GCC TTG GCC CTT CGT
725
Phe Gly Val Asp Pro Ser Ala Asp Pro Glu Leu Alo Leu Ala Leu Arg>
210 215 220

GTA TCT ATG GAA GAG CAG CGG CAC GCA GGA GGA GGA GCG CGG CGG GCA
773
Val Ser Met Glu Glu Gln Arg His Ala Gly Gly Gly Ala Arg Arg Ala>
225 230 235

GCT CGA GCT TCT GCT GCT GAG GCC GGG ATT GCT ACG ACT GGG ACT GAA
821
Ala Arg Ala Ser Ala Ala Glu Ala Gly Ila Ala Thr Thr Gly Thr Glu>
240 245 250

GAC TCA GAC GAT GCC CTG CTG AAG ATG ACC ATC AGC CAG CAA GAG TTT
869
Asp Ser Asp Asp Ala Leu Leu Lys Met Thr Ile Ser Gln Gln Glu Phe>
255 260 265

GGC CGC ACT GGG CTT CCT GAC CTA AGC AGT ATG ACT GAG GAA GAG CAG
917
Gly Arg Thr Gly Leu Pro Asp Leu Ser Ser Met Thr Glu Glu Glu Gln>
270 275 280 285

ATT GCT TAT GCC ATG CAG ATG TCC CTG CAG GGA GCA GAG TTT GGC CAG
965
Ile Ala Tyr Ala Met Gln Met Ser Leu Gln Gly Ala Glu Phe Gly Gln>
290 295 300

GCG GAA TCA GCA GAC ATT GAT GCC AGC TCA GCT ATG CAC ACA TCT GAG
1013
Ala Glu Ser Ala Asp Ile Asp Ala Ser Ser Ala Met Asp Thr Ser Glu>
305 310 315

CCA GCC AAG GAC GAG GAT GAT TAC GAC CTG ATG CAG GAC CCC GAG TTC
1061
Pro Ala Lys Glu Glu Asp Asp Tyr Lsp Vol Met Gln Asp Pro Glu Phe>
320 325 330

CTT CAG AGT GTC CTA GAG AAC CTC CCA GGT GTG GAT CCC AAC AAT GAA
1109
Leu Gln Ser Val Leu Glu Asn Leu Pro Gly Val Asp Pro Asn Asn Glu>
335 340 345

GCC ATT CGA AAT GCT ATG GGC TCC CTG CCT CCC AGG CCA CCA AGG ACG
1157
Ala Ile Arg Asn Ala Met Gly Ser Leu Pro Pro Arg Pro Pro Arg Thr>
350 355 360 365

GCA AGA AGG ACA ACA AGG AGG AAG ACA AGA AGT CAG ACT GGA GGG AAA
1285
Alo Arg Arg Thr Arg Arg Arg Lys Thr Arg Ser Glu Thr Gly Gly Lys>
370 375 380

GGG
TAGCTGAGTCTGCTTAGGGGACTGCATGGGAACCACGGAATATAGGGTTAGATGTGTG
T
Gly >
TATCTGTAACCATTACAGCCTAAATAAAGCTTGGCAACTTTTAAAAAAAAAAAAAAAA
A